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10/584,002	06/22/2006	Mark Derek Cregan	07-2353	6304
20306 7590 04/28/2008 MCDONNELL BOEHNEN HULBERT & BERGHOFF LLP			EXAMINER	
300 S. WACK	ER DRIVE	The Beneficial Lies	SAJJADI, FERE	YDOUN GHOTB
32ND FLOOR CHICAGO, IL			ART UNIT	PAPER NUMBER
,			1633	
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			04/28/2008	PAPER

Please find below and/or attached an Office communication concerning this application or proceeding.

The time period for reply, if any, is set in the attached communication.

Office Action Summary

Application No.	Applicant(s)	
10/584,002	CREGAN ET AL.	
Examiner	Art Unit	
FEREYDOUN G. SAJJADI	1633	

- The MAILING DATE of this communication appears on the cover sheet with the correspondence address — Period for Reply A SHORTENED STATUTORY PERIOD FOR REPLY IS SET TO EXPIRE 1 MONTH(S) OR THIRTY (30) DAYS, WHICHEVER IS LONGER, FROM THE MAILING DATE OF THIS COMMUNICATION. - Extension of some may be available under the provisions of 37 CFR 1.736(s), in no event, however, may a reply be limited in the provision of the provision of 17 CFR 1.736(s), in no event, however, may a reply be limited in the provision of 17 CFR 1.736(s), in no event, tho event, however, may a reply be limited in the provision of 17 CFR 1.736(s), in no event, however, may a reply be limited in the communication of the provision of 17 CFR 1.736(s), in no event, however, may a reply be limited in the communication. - If all the to reply within the set or ordended period for reply with p. Office lates than throe months after the mailing date of this communication, even if timely filled, may reduce any earned patient term adjustment. See 37 CFR 1.704(s). Status 1) Responsive to communication(s) filled on 12 July 2007. 2a) This action is FINAL. 2b) This action is non-final. 3) Since this application is in condition for allowance except for formal matters, prosecution as to the merits is closed in accordance with the practice under Ex parte Quayle, 1935 C.D. 11, 453 O.G. 213. Disposition of Claims 4) Claim(s) 1-24 is/are pending in the application. 4a) Of the above claim(s) is/are allowed. 5) Claim(s) is/are allowed. 5) Claim(s) is/are allowed. 7) Claim(s) is/are objected to. 8) Claim(s) is/are rejected. 7) Claim(s) is/are rejected. 7) Claim(s) is/are rejected. 7) The specification is objected to by the Examiner. Applicant may not request that any objection to the drawing(s) be held in abeyance. See 37 CFR 1.85(a). Replacement drawing sheet(s) including the correction is required if the drawing(s) is objected to. See 37 CFR 1.121(d).
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11) I he oath or declaration is objected to by the Examiner. Note the attached Office Action or form PTO-152.
Priority under 35 U.S.C. § 119
12) Acknowledgment is made of a claim for foreign priority under 35 U.S.C. § 119(a)-(d) or (f).
a)
 Certified copies of the priority documents have been received.
Certified copies of the priority documents have been received in Application No
 Copies of the certified copies of the priority documents have been received in this National Stage
application from the International Bureau (PCT Rule 17.2(a)).
* See the attached detailed Office action for a list of the certified copies not received.
Attachment(s)
1) Notice of References Cited (PTO-892) 4) Interview Summary (PTO-413)
2) Notice of Draftsperson's Patent Drawing Review (PTO-948) 3) Information Disclosure Statement(s) (PTO/SS/CE) 7) Notice of Informal Patent Application

Paper No(s)/Mail Date _____.

6) Other: _____.

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DETAILED ACTION

Claims 1-30 are pending in the Application,

Election/Restrictions

Restriction is required under 35 U.S.C. 121 and 372.

 This application contains the following inventions or groups of inventions which are not so linked as to form a single general inventive concept under PCT Rule 13.1.

In accordance with 37 CFR 1.499, applicant is required, in reply to this action, to elect a single invention to which the claims must be restricted.

Group I, claim(s) 1-18, drawn to a method for isolating progenitor cells having stem-cell-like characteristics from human mammary secretions.

Group II, claim(s) 19-24, drawn to a method of creating cells or tissues in a mother or infant comprising administering to said mother or infant pluripotent or multipotent progenitor cells.

Please note that PCT Rule 13.2, no longer specifies the combinations of categories of invention which are considered to have unity of invention. The categories of invention in former PCT Rule 13.2 have been replaced with a statement describing the method for determining whether the requirement of unity of invention is satisfied. Unity of invention exists only when there is a technical relationship among the claimed inventions involving one or more special technical features. The term "special technical features" is defined as meaning those technical features that define a contribution which each of the inventions considered as a whole, makes over the prior art.

The inventions listed as Groups I and II do not relate to a single general inventive concept under PCT Rule 13.1 because, under PCT Rule 13.2, they lack the same or corresponding special technical features for the following reasons:

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According to PCT Rule 13.2, unity of invention exists only when a shared same or corresponding special technical feature is a contribution over the prior art. The technical feature, which is shared by Groups I and II, is a stem-cell-like progenitor cell isolated from mammary secretions.

Groups I and II do not share a special technical feature over the art because the inventions lack an inventive step under PCT Article 33(3) as being obvious over Young et al. (Aus. J. Zool. 45(4):423-433; 1997), who describe the isolation of vacuolated mononuclear cells from milk, closely resembling blast cells that may be primitive stem cells. (Abstract and p. 425).

The claims in Groups I and II are drawn to distinct methods, directed to separate goals, employing distinct method steps, requiring non-coextensive search and examination. Thus, it follows from the preceding analysis that the claimed inventions listed as Groups I and II do not relate to a single inventive concept under PCT Rule 13.1 because, under PCT Rule 13.2, they lack the same or corresponding technical features for the reasons set forth above.

This application contains claims directed to more than one species of the generic
invention. These species are deemed to lack unity of invention because they are not so linked as
to form a single general inventive concept under PCT Rule 13.1.

The species are as follows:

A specifically named single species of human, either a male or a female, as recited in claim 1.

A specifically named single species of mammary secretion, either colostrum, mature milk or dry period secretion, as recited in claim 1.

A specifically named single species for the period of secretion, either non-pregnant, pregnant, lactating or involuting period, as recited in claims 1.

A specifically named single species enzyme, either DNAse, Proteinase or RNase, as recited in claim 10

A specifically named single species as the target of cell therapy, either a mother or an infant, as recited in claim 19.

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A specifically named single species of therapy, either gene therapy or intrauterine fetal treatment, as recited in claim 21.

A specifically named single species for the type of therapy, either bioengineering, lactoengineering, breast tissue regeneration, breast reconstructive surgery, or cosmetic or enhancement surgery, exocrine gland tissue regeneration or its combination with surgery, as recited in claim 24

Applicant is required, in reply to this action, to elect a single species to which the claims shall be restricted if no generic claim is finally held to be allowable. The reply must also identify the claims readable on the elected species, including any claims subsequently added. An argument that a claim is allowable or that all claims are generic is considered non-responsive unless accompanied by an election.

Upon the allowance of a generic claim, applicant will be entitled to consideration of claims to additional species which are written in dependent form or otherwise include all the limitations of an allowed generic claim as provided by 37 CFR 1.141. If claims are added after the election, applicant must indicate which are readable upon the elected species. MPEP § 809.02(a).

The claims are deemed to correspond to the species listed above in the following manner:

Claims 1, 10, 19, 21, 24 and claims dependent therefrom correspond to all the species listed above.

The following claim(s) are generic: 1-24.

The species listed above do not relate to a single general inventive concept under PCT Rule 13.1 because, under PCT Rule 13.2, the species lack the same or corresponding special technical features for the following reasons:

The species listed above do not relate to a single general inventive concept under PCT Rule 13.1 because, under PCT Rule 13.2, the species lack the same or corresponding special technical features for the following reasons: under PCT Rule 13.2, the species lack the same or corresponding special technical features for the following reasons: As the technical features for

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colostrum, mature milk or dry period secretion, non-pregnant, pregnant, lactating or involuting periods, DNAse, Proteinase or RNase, cell therapy, gene therapy as maternal or infant, bioengineering, lactoengineering, breast tissue regeneration, breast reconstructive surgery, or cosmetic or enhancement surgery, exocrine gland tissue regeneration or its combination with surgery, linking the members do not constitute a special technical feature as defined by PCT Rule 13.2, particularly since each of the species is structurally and likely functionally distinct, or requires non-commonly shared particulars, is capable of separate utility, and does not share a substantially common structural feature, the requirement for unity of invention is not fulfilled.

Applicant is advised that the reply to this requirement to be complete must include (i) an election of a species or invention to be examined even though the requirement may be traversed (37 CFR 1.143) and (ii) identification of the claims encompassing the elected invention.

The election of an invention or species may be made with or without traverse. To preserve a right to petition, the election must be made with traverse. If the reply does not distinctly and specifically point out supposed errors in the restriction requirement, the election shall be treated as an election without traverse.

Applicant is reminded that upon the cancellation of claims to a non-elected invention, the inventorship must be amended in compliance with 37 CFR 1.48(b) if one or more of the currently named inventors is no longer an inventor of at least one claim remaining in the application. Any amendment of inventorship must be accompanied by a request under 37 CFR 1.48(b) and by the fee required under 37 CFR 1.17(i).

Any inquiry concerning this communication or earlier communications from the examiner should be directed to FEREYDOUN G. SAJJADI whose telephone number is (571)272-3311. The examiner can normally be reached on 6:30 AM-3:30 PM EST.

If attempts to reach the examiner by telephone are unsuccessful, the examiner's supervisor, Joseph Woitach can be reached on (571) 272-0739. The fax phone number for the organization where this application or proceeding is assigned is 571-273-8300.

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Information regarding the status of an application may be obtained from the Patent Application Information Retrieval (PAIR) system. Status information for published applications may be obtained from either Private PAIR or Public PAIR. Status information for unpublished applications is available through Private PAIR only. For more information about the PAIR system, see http://pair-direct.uspto.gov. Should you have questions on access to the Private PAIR system, contact the Electronic Business Center (EBC) at 866-217-9197 (toll-free). If you would like assistance from a USPTO Customer Service Representative or access to the automated information system, call 800-786-9199 (IN USA OR CANADA) or 571-272-1000.

/Fereydoun G Sajjadi/

Fereydoun G. Sajjadi, Ph.D. Examiner, Art Unit 1633